SAFEGUARDS:

Information contained in the system is maintained in accordance with DFBP procedures. Manual information in the system is safeguarded in locked file cabinets within a limited access room in a limited access building. Access to manual files is limited to personnel who have a need for files to perform official duties. Operational access to information maintained on a dedicated computer system, is controlled by levels of security provided by password keys to prevent unauthorized entry, and an audit trail of accessed information. Access is also limited to personnel who have a need to know to perform official duties.

RETENTION AND DISPOSAL:

Data is maintained for current and prior years in a master file. Data is not destroyed, but maintained for historical purposes.

SYSTEM MANAGER(S) AND ADDRESS:

Director, DFBP, Office of Justice Programs, 810 Seventh Street, NW, Washington, DC 20531.

NOTIFICATION PROCEDURE:

Same as above.

RECORD ACCESS PROCEDURES:

A request for access to a record from the system shall be in writing, with the envelope and letter marked ''Privacy Access Request.'' Direct the access request to the System Manager listed above. Identification of individuals requesting access to their records will include fingerprinting (28 CFR 20.34).

CONTESTING RECORDS PROCEDURES:

An individual desiring to contest or amend information maintained in the system should direct the request to the System Manager listed above. The request should state clearly and concisely the information being contested, the reasons for contesting the information, and the proposed information amendment(s) sought.

RECORD SOURCE CATEGORIES:

Sources of information contained in the system are Federal and State courts, individuals convicted of certain drug offenses, individuals convicted of defense-contract related felonies, United States Attorneys, and Federal agencies.

SYSTEMS EXEMPTED FROM CERTAIN PROVISIONS OF THE ACT:

None.

[FR Doc. 99–11662 Filed 5–7–99; 8:45 am]

BILLING CODE 4410–CJ–M

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

Manufacturer of Controlled Substances; Notice of Registration

By Notice dated January 27, 1999, and published in the **Federal Register** on February 10, 1999, (64 FR 6684), Isotec, Inc., 3858 Benner Road, Miamisburg, Ohio 45342, made application by renewal to the Drug Enforcement Administration (DEA) to be registered as a bulk manufacturer of the basic classes of controlled substances listed below:

Drug	Schedule
Cathinone (1235)	I
Methcathinone (1237)	I
N-Ethylamphetamine (1475)	I
N,N-Dimethylamphetamine (1480)	I
Aminorex (1585)	I
Methaqualone (2565)	I
Lysergic acid diethylamide (7315)	I
Tetrahydrocannabinols (7370)	I
Mescaline (7381)	I
2,5-Dimethoxyamphetamine (7396).	I
3,4-Methylenedioxyamphetamine (7400).	I
3,4-Methylenedioxy-N-ethylamphetamine (7404).	I
3,4-Methylenedioxymethamphetamine (7405).	I
4-Methoxyamphetamine (7411)	I
Psilocybin (7437)	I
Psilocyn (7438)	I
N-Ethyl-1-phenylcyclohexylamine (7455).	I
Dihydromorphine (9145)	I
Normorphine (9313)	I
Acetylmethadol (9601)	I
Alphacetylmethadol Except Levo-Alphacetylmethadol (9603).	I
Normethadone (9635)	I
3-Methylfentanyl (9813)	I
Amphetamine (1100)	II
Methamphetamine (1105)	II
Methylphenidate (1724)	II
Amobarbital (2125)	II
Pentobarbital (2270)	II
Secobarbital (2315)	II
1-Phenylcyclohexylamine (7460)	II
Phencyclidine (7471)	II
Phenylacetone (8501)	II
1-Piperidinocyclohexanecarbonitrile (8603).	II
Codeine (9050)	II
Dihydrocodeine (9120)	II
Oxycodone (9143)	II
Hydromorphone (9150)	II
Benzoylecgoning (9180)	II
Ethylmorphine (9190)	II
Hydrocodone (9193)	II
Isomethadone (9226)	II
Meperidine (9230)	II
Methadone (9250)	II
Methadone intermediate (9254)	II
Dextropropoxyphene, bulk (non-dosage forms) (9273).	II
Morphine (9300)	II
Levo-Alphacetylmethadol (9648)	II
Oxymorphone (9652)	II
Fentanyl (9801)	II

The firm plans to use small quantities of the listed controlled substances to produce standards for analytical laboratories.

DEA has considered the factors in Title 21, United States Code, Section 823(a) and determined that the registration of Isotec, Inc. to manufacture the listed controlled substances is consistent with the public interest at this time. DEA has investigated Isotec, Inc. on a regular basis to ensure that the company's continued registration is consistent with the public interest. These investigations have included inspection and testing of the company's physical security systems, audits of the company's records, verification of the company's compliance with state and local laws, and a review of the company's background and history. Therefore, pursuant to 21 U.S.C. 823 and 28 CFR 0.100 and 0.104, the Deputy Assistant Administrator, Office of Diversion Control, hereby orders that the application submitted by the above firm for registration as a bulk manufacturer of the basic classes of controlled substances listed above is granted.

Dated: April 26, 1999.

John H. King,

Deputy Assistant Administrator, Office of Diversion Control, Drug Enforcement Administration.

[FR Doc. 99–11693 Filed 5–7–99; 8:45 am]

BILLING CODE 4410–09–M

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[Docket No. 96–41]

Paul W. Saxton, Continuation of Registration

On July 15, 1996, the Deputy Assistant Administrator, Office of Diversion Control, Drug Enforcement Administration (DEA) issued an Order to Show Cause to Paul W. Saxton, D.O. (Respondent) of Sandy, Utah, notifying him or an opportunity to show cause as to why DEA should not revoke his DEA Certificate of Registration AS9420059 and deny any pending applications for renewal of such registration as a practitioner pursuant to 21 U.S.C. 823(f) and 824(a)(4), for reason that his continued registration would be inconsistent with the public interest.